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- (71) Applicant (for all designated States except US): PHARMNSEAS, INC. [US/US]; 22512 Avenida Empresa, Rancho Santa Margarita, CA 92688 (US).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): HOWARD, Larry [US/US]; 25262 Arcadian Avenue, Mission Viejo, CA 92691 (US).

- (74) Agent: CONNORS, John, J.; Connors & Associates, Suite 220, 1600 Dove Street, Newport Beach, CA 92660-2427 (US).
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1/26646 A

(54) Title: NUTRACEUTICAL PRODUCTS CONTAINING SAMe AND DIETARY SUPPLEMENTS AND METHOD OF MAN-UFACTURE AND USE THEREOF

(57) Abstract: A nutraceutical product comprises a mixture SAMe and a dietary supplement, where the moisture content of the product is less than 5 weight percent. A nutraceutical preferred product includes a mixture of (RS)-(+)-SAMe and (SS)-(+)-SAMe diastereoisomers, with the (SS)-(+)-SAMe diastereoisomer being at a concentration of at least 95 weight percent of the mixture.

NUTRACEUTICAL PRODUCTS CONTAINING SAME AND DIETARY SUPPLEMENTS & METHOD OF MANUFACTURING AND USE THEREOF

RELATED APPLICATIONS

This utility application is based on the following United States provisional patent applications: Serial No. 60/158,298, filed October 8, 1999, entitled Time Release Coated Natural Supplement; Serial No. 60/158,328, filed October 8, 1999, entitled Herbal Mind And Mood Support Composition; Serial No. 60/158,329, filed October 8, 1999, entitled Herbal Stress Support Composition; Serial No. 60/158,480, filed October 8, 1999, entitled Joint Support Composition; and Serial No. 60/158,482, filed October 8, 1999, entitled Herbal Diet And Energy Support Composition. All of these provisional patent applications are incorporated herein by reference and made a part of this application.

BACKGROUND OF THE INVENTION

S-adenosyl-L-methionine, and its salts, (either or both herein referred to as SAMe) are well known pharmacologically active compositions that combat depression, arthritis, and liver diseases such as, for example, cirrhosis. SAMe occurs as two diastereoisomers: (RS)-(+)-SAMe and (SS)-(+)-SAMe. The (SS)-(+)-SAM-e diastereoisomer is the pharmacologically active diastereoisomer, and SAMe products containing (SS)-(+)-SAMe at a concentration of at least 95 weight percent (%) of the total diastereoisomers mixture are known. A suitable source of SAMe, including the SAMe with the higher concentration of the (SS)-(+)-SAMe diastereoisomer, is Gnosis S. r. l. of Milan, Italy.

SUMMARY OF THE INVENTION

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This invention is a nutraceutical product comprising a blend of SAMe and one or more dietary supplements. This nutraceutical product may be used by both humans and animals. Different nutraceutical products are provided having multiple beneficial properties. In this invention the dietary supplements enhance the beneficial effects of the SAMe, or provide additional beneficial effects, or both. The nutraceutical products of this invention preferably employ SAMe with the higher concentration of the (SS)-(+)-SAMe diastereoisomer.

The nutraceutical products of this invention may be administered by any conventional route such as, for example, oral, rectal, nasal, topical, or pareateral tablets, caplets, softgel pills, capsules, as suspensions, emulsions, solutions, suppositories, sprays, gums, drops, lozenges, or injectibles They may be sold either in bulk to companies that repackage the product for the consumer or in a form suitable for use directly by the consumer. For example, if ingested orally, the nutraceutical products of this invention are in the form of a softgel pill, caplet, capsule or tablet. Preferably, the softgel pill, caplet, capsule or tablet is treated with a time release agent. An enteric coating and/or micro-encapsulation may be employed to achieve time release, and one suitable time release agent is an aqueous ethyl cellulose dispersion sold under the trademark Surelease® by Colorcon, Inc. of Santa Ana, California.

Packaging of the nutraceutical products of this invention is important. As discussed subsequently in greater detail, moisture adversely affects SAMe. Retailers of nutraceutical products usually conduct accelerated aging tests by subjecting the packaged product to high temperatures and high humidity over a controlled time duration, measuring the SAMe concentration in the product before and after

testing. It has been discovered that the softgel pill, caplet, capsule or 1 2 tablet form of the nutraceutical products of this invention completely 3 enclosed and sealed within an aluminum foil package resist degradation. Preferably, the softgel pill, caplet, capsule or tablet is first completely 4 5 enclosed within a plastic wrap and then completely enclosed and sealed 6 the aluminum foil. The aluminum foil package acts as a moisture barrier, enabling said softgel pill, caplet, capsule or table to have 7 substantially prolonged life when subjected to accelerated testing than a 8 softgel pill, caplet, capsule or table contained within, for example, a 9 sealed bottle or plastic blister pack. Preferably, the thickness of the 10 11 aluminum foil is from 2 to 6 mils.

The nutraceutical products of this invention may also be sold in forms other than a softgel pill, caplet, capsule or tablet. For example these products may be used as topical applications such as creams, gels, patches, etc. They may also be in liquid form especially when used as an injectibles, or even as an enemas.

In the preferred embodiments of this invention the following dietary supplements are used:

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Co-Q-10 (Coenzyme Q-10), 20

> St. John's Wort derivative, either powder or any form of extract, Ginkgo Biloba derivative, either powder or any form of extract, Ma Huang derivative, either powder or any form of extract,

Epimeduim pinnatum derivative, either powder or extract,

Guarana derivative, either leaf or root powder or any form of 25

26 extract.

0126646A1 L >

Citrus Aurantium derivative, either powder or any from of extract, 27 28

Kava Kava derivative, either powder or any form of extract,

cetyl myristoleate (CMO),

cetyl myristoleate blended with cetyl oleate, cetyl esters and cetyl myristate,

- 3 Valerian derivative, either powder or any form of extract,
- 4 glutathione,
- fructose 1,6 diphosphate (FDP),
- 6 glucosamine sulfate, glucosamine HCL, or glucosamine potassium,
- methyl sulfanyl methane (MSM),
- 8 melatonin,
- 9 hydrolyzed collagen,
- 10 chondroitin sulfate,
- 11 citicoline,
- 12 alpha GPC.

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Any one, or combinations of two or more, of these dietary supplements are mixed with SAMe, preferably with the SAMe comprising from about 0.1 to about 99 weight percent and the dietary supplement comprising from about 1 to about 99 weight percent of the nutraceutical product.

SAMe in decomposes the presence of water and high This presents a problem when blending with dietary temperatures. In accordance with this invention, during manufacture of supplements. the nutraceutical products of this invention conditions are controlled so that the moisture content of the product is less than 5 weight percent, preferably less than 3 weight percent. In some instances this may require drying the dietary supplements prior to blending with the SAMe. In the preferred manufacturing method the SAMe and dietary supplement or supplements being mixed with the SAMe are dry powders having a mesh ranging from about 60 to about 90. The humidity of the manufacturing environment of nutraceutical product is less than 25% and the temperature of the manufacturing environment of nutraceutical product is from 20 to 15 degrees Celsius. The

nutraceutical product is encapsulated within capsules within 2 hours after manufacture. Preferably, these capsules are enteric coated, or otherwise treat with a time release agent, within 2 hours after encapsulation.

This invention also includes a method of supplementing a diet by administering for at least 10 to 14 days SAMe at a dosage of from 800 to 1200 milligrams per day and at the same time administering a nutraceutical product comprising SAMe and a dietary supplement, said nutraceutical product including a a dosage of at least 400 milligrams per day of SAMe, and after 14 days administering only said nutraceutical product.

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DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

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STRESS REDUCTION COMPOSITION

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The preferred stress reduction compositions of this invention preferably include a combination of SAMe and other natural stress reducing substances in effective amounts for maintaining and/or improving emotional and physical health during periods of stress. One preferred embodiment of this invention especially effective in reducing emotional and physical stress and improving the ability to cope with such stress is SAMe in combination with one or more of the following: a Kava Kava derivative and a Valerian derivative.

The following are the preferred weight percentage ranges of these stress reduction compositions:

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28 Component Weight Percentage

29 30

31 Kava Kava 1-70% Extract Activity

about 5% to about 90%

about 5% to about 90%

SAMe

Valerian 1-30% Extract Activity

about 5% to about 90%

For efficacy in providing stress reduction, SAMe is administered in dosages ranging from 10 mg to 2000 mg per day. Most preferably, SAMe is present in an amount of 400 mg per total daily dose which is to be taken one to two times per day. Kava Kava extract is derived from a tropical plant found in the tropical regions of the Pacific rim and is known and used for its benefit of relieving symptoms of stress and anxiety, and to promote and induce relaxation. When used in combination with SAMe, a Kava Kava extract, preferably 1-70% by weight active ingredient, is present in an amount ranging from 10 mg to 2000 mg per dose per day. Preferably, a 30% by weight active ingredient Kava Kava extract is present in an amount of 600 mg per total daily dose taken one to two times.

Another composition which is effective in providing health benefits in the form of stress reduction and/or coping is Valerian Extract. Valerian is an herbal product known for its anti-anxiety and stress benefits and is sometimes used to help promote sleep. When used in combination with SAMe, a Valerian Root extract, preferably .8-30% by weight active ingredient is present in an amount ranging from 10 mg to 2000 mg per dose per day. Preferably, a 0.8% by weight Valerian Root extract, is present in an amount of 100 mg per total daily dose taken one to two times per day.

Another preferred composition is to combine all of the above: 200 mg SAMe with 600 mg Kava Kava extract, and 100 mg Valerian root extract within a time-release or enteric coated capsule, caplet, or tablet along with standard inert ingredients utilized in the art. This composition to be taken one to two times per day.

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	WO 01/26646 PC1/US00/2/559
1	Preferred Dosages
2 3 4 5	 For the first ten days: Take 800 - 1,200 mg per day of SAMe Active Ions- in addition to the Stress Support product.
6 7 8	2. Afterwards, continue only taking the Stress Support product daily.
9 10 11	 400 mg of SAMe 600 mg of Kava Kava Extract, 30-50% Kavalactones 100 mg Valerian Root Extract (.8 - 1%) Valerenic acid
13	WEIGHT REDUCTION AND ENERGY
14	ENHANCEMENT COMPOSITION
15 16	The preferred weight reduction and energy enhancement
17	compositions of this invention preferably include a combination of
18	SAMe and one or more other natural appetite suppressant and/or
19	energy enhancing substances in effective amounts for this purpose. One
20	preferred embodiment of this invention especially effective for weight
21	control and energy enhancement is SAMe in combination with one or
22	more of the following: a Ma Huang derivative, a Epimedium Pinnatum
23	derivative, a Citrus Aurantium derivative, and a Guarana derivative.
24	These substances suppress appetite and increase energy. These
25	substances provide weight reduction benefits by speeding up the body's
26	metabolism, thus increasing the body's ability to burn fat without
27	exercise and enhance energy levels.
28	The following are the preferred weight percentage ranges of these
29	weight reduction and energy enhancement compositions:
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31	Component Weight Percentage
32	SAMe about 5% to about 90%
33	Ma Huang Extract 1-8% Activity about 5% to about 90%

about 5% to about 90%

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Epimedium Extract 1-50% Activity

Citrus Aruantium Extract 1-40% Activity 1 about 5% to about 90%

2 Guarana Extract 1-50% Activity about 5% to about 90%

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For efficacy in providing weight loss and energy benefits, SAMe is present in an amount ranging from 10 mg to 2000 mg per day. Preferably SAMe is present in an amount of 400 mg per day two times Other weight loss benefit compositions may be taken with SAMe.

Another composition that is effective in providing health benefits in the form of weight reduction is Ma Huang. Ma Huang is a naturally occurring herb which contains as an active weight reduction ingredient sympathomimetic compound Ephedra Sinica, or its extract ephedrine. When used in combination with SAMe, Ma Huang Extract, preferably at a concentration of 1-8% by weight Activity, is present in an amount ranging from 10 mg to 2000 mg per dose per day. Preferably, a 6% by weight Ma Huang extract is present in an amount of 300 mg per daily dose taken two times per day before morning and evening meal.

composition which is effective in providing Another benefits in the form of weight reduction is Epimedium Pinnatum Extract. Epimedium Extract is known to increase energy levels and metabolism. When used in combination with SAMe, Epimedium Extract, preferably at a concentration of 1-50% by weight Activity is present in an amount ranging from 10 mg to 2000 mg per dose. Preferably, a 10% by weight Epimedium Pinnatum extract is present in an amount of 100 mg per daily dose taken two times per day before morning and evening meal.

composition which is effective in providing Another benefits in the form of weight reduction is Citrus Aurantium Extract. Citrus Aurantium Extract is known to increase energy levels and metabolism. When used in combination with SAMe, a 1-40% by weight

Activity concentration of Citrus Aurantium Extract is present in an amount ranging from 10 mg to 2000 mg per daily dose. Preferably, a 4% by weight Citrus Aurandum extract is present in an amount of 100 mg per dose taken two times per day before morning and evening meal.

Another composition which is effective in providing health benefits in the form of weight reduction is Guarana Extract. Guarana Extract is known to increase energy levels and metabolism. When used in combination with SAMe, a 1-50% by weight Activity concentration of Guarana is present in an amount ranging from 10 mg to 2000 mg daily per dose. Preferably, a 10% by weight caffeine Guarana extract is present in an amount of 80 mg per dose taken two times per day.

Another preferred composition is to combine all of the above for an average daily dosage as follows: 200 mg to 400 mg SAMe with 900 mg Ma Huang Extract, 200 mg Epimedium Pinnatum Extract, 100 mg Citrus Aurantium Extract and 100 mg Guarana Extract within a time-release coated capsule or tablet along with standard inert ingredients utilized in the art.

Another preferred composition is to combine all of the above: 400 mg SAMe with 300 mg Ma Huang Extract, 100 mg Epimedium Pinnatum Extract, 100 mg Citrus Aurantium Extract and 80mg Guarana Extract within a time-release coated capsule or tablet along with standard inert ingredients utilized in the art; said composition to be taken two times per day before morning and evening meal.

Preferred Dosages

1. For the first ten days: Take 800 - 1,200 mg per day of SAMe 400 mg Active Ions tablets, caplets, or capsules — in addition to our Diet Support

2. Afterwards, continue only taking the Diet Support composition daily.

•	400 mg of Active SAMe Ions
•	300-400 mg of Ma Huang Extract 6% Alkaloids
•	100 mg Epimedium Pinnatum Extract 10-30% Icariin
•	100-200 mg Citrus Aurantium Extract, 4-6% Synephrine
•	80-120 mg Guarana Seed Extract 10% Caffeine

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JOINT ANTI-INFLAMMATION COMPOSITION

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The preferred anti-inflammation compositions of this invention preferably include a combination of SAMe and other natural inflammation substances in effective amounts for this purpose. embodiment of this invention especially effective inflammation reduction and joint pain is SAMe in combination with one or more of the following: methyl sulfanyl methane, cetyl myristoleate, glucosamine sulfate, glucosamine HCL, or glucosamine potassium, and hydrolyzed collagen. These components are known to help (a) relieve symptoms of arthritis and inflammation, (b) rebuild connective tissue, and (c) relieve symptoms related to sport injury and accidents affecting joints. There is a benefit to be found in taking these substances in combination, which increases the overall health benefit to joints over that obtained by taking each substance individually.

The following are the preferred weight percentage ranges of these anti-inflammation compositions:

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26	Component	Weight Percentage
27	SAMe	about 5% to about 90%
28	methyl sulfanyl methane	about 5% to about 90%
29	Cetyl myristoleate 1-99% Pure	about 5% to about 90%
30	Glucosamine sulfate, HCL or potassium	about 5% to about 90%
31	Hydrolyzed collagen	about 5% to about 90%

For efficacy in providing joint health benefits, SAMe is administered in an amount ranging from 10 mg to 2000 mg per day.

3 Preferably SAMe is present in an amount ranging from 400 mg to 800

mg per day. Most preferably, SAMe is present in an amount of 100 mg

per dose which is to be taken two to three times per day. Other joint

benefit compositions may be taken with SAMe.

Another composition which is effective in providing health benefits to joints, is MSM (methyl sulfanyl methane). MSM is known to reduce symptoms caused by arthritis and sport injuries. When used in combination with SAMe, MSM is present in an amount ranging from 10 mg to 5000 mg per daily dose. Preferably, MSM is present in an amount of 300 mg per dose taken two times per day.

Another composition which is effective in providing health benefits to joints, is cetyl myristoleate. Cetyl mysristoleate is known to help symptoms of rhumotoid arthritis and inflammation. When used in combination with SAMe, a 1-99% by weight concentration of cetyl myristoleate is present in an amount ranging from 0.5 mg to 2000 mg per dose per day. Preferably, an 20% by weight concentration of cetyl mysristoleate is present in an amount of 50 mg per dose taken two times per day.

Another composition which is effective in providing health benefits to joints is glucosamine sulfate, glucosamine HCL, or glucosamine potassium. This composition is known to help rebuild connective tissues and to ease symptoms of arthritis, injury, and inflammation. When used in combination with SAMe, glucosamine sulfate, glucosamine HCL, or glucosamine potassium is present in an amount ranging from 10 mg to 2000 mg per dose per day. Preferably, glucosamine sulfate, glucosamine HCL, or glucosamine potassium is present in an amount of 1000 mg per dose taken two to three times per day.

Another preferred composition is to combine all of the above: 100 mg SAMe with 300 mg MSM, 50 mg CMO, and 1000 mg glucosamine sulfate within a time-release coated capsule or tablet along with standard inert ingredients utilized in the art; said composition to be taken one to four times per day.

Another preferred composition is to combine all of the above: about 400 mg SAMe with about 1000 mg MSM, with about 75 mg cetyl myristoleate, with about 1,000 mg glucosamine sulfate, with about 100 mg hydrolyzed collagen within a time-release coated capsule or tablet.

Preferred Dosages

1. For the first ten days: Take 800 - 1,200 mg per day of SAMe 400mg Active Ions tablets, caplets, or capsules - in addition to the Joint Support composition

2. Afterwards, continue only the Joint Support daily.

• 400 mg of Active SAMe Ions

• 1000 - 1,500 mg of Glucosamine Sulfate (or HCL or Potassium)

• 300-500 mg of MSM (Methylsulfonylmethane)

 • 50-500 mg of CMO blend, 20% Cetyl Myristoleate

MIND AND MOOD ENHANCEMENT COMPOSITION

The preferred mind and mood enhancement compositions of this invention preferably include a combination of SAMe and other natural mind and mood substances in effective amounts for this purpose. One preferred embodiment of this invention especially effective for mind and mood enhancement is SAMe in combination with one or more of the following: a St. John's Wort derivative and a Ginko Biloba derivative. These emotional balance and brain function compositions are presented in a variety of formulations, with or without other emotional balance

1 and/or brain function ingredients.

According to the invention, the herbal treatment composition preferably includes the desired combination of natural mood and brain function promoting substances in effective amounts for maintaining and/or improving emotional balance and/or memory and/or function. The foregoing descriptions of quantity and dosage are intended to be specific examples and are not intended to limit the scope of the invention.

The following are the preferred weight percentage ranges of these mind and mood enhancement compositions:

12	Component	Weight Percentage
13	SAMe	about 5% to about 90%
14	St. John's Wort .1-50%	about 5% to about 90%
15	Ginkgo Biloba 1-99%	about 5% to about 90%

For efficacy in providing mind and mood benefits, SAMe is administered in an amount ranging from 10 mg to 2000 mg per day. Preferably, SAMe is used in an amount 400 mg per day. Most preferably, SAMe is used in an amount of 400 mg per daily dose which is to be taken one to two times per day.

Another composition which is effective in providing health benefits in the form of increased emotional balance is St. John's Wort. St. John's Wort is a naturally occurring herb which contains the therapeutically active ingredient hypericin and hyperforin. These ingredients have an anti-depressive effect which is believed to be caused by modifying serotonin levels in the brain. When used in combination with SAMe, St John's Wort, preferably a concentration between .1-50% by weight, is used in an amount ranging from 10 mg to 5000 mg per dose per day. Preferably a 0.3% by weight hypericin

1 concentration of St. John's Wort extract is used in an amount of 900 mg 2 per daily dose taken two times per day.

Another composition that is effective in providing health benefits in the form of improving memory and brain function as well as acting as an anti-depressant is Ginko Biloba. When used in combination with SAMe, Ginko Biloba, preferably a concentration between 1-99% by weight, is used in an amount ranging from 0.5 mg to 2000 mg per dose per day. Preferably, a 24/6% by weight Ginko Biloba extract, is used in an amount of 60 mg per daily dose.

Another preferred composition is to combine all of the above: 200 to 400 mg SAMe with 60 mg Ginko Biloba Extract, and 900 mg St. John's Wort extract within a time-released coated capsule or tablet along with standard inert ingredients utilized in the art; said composition to be taken one to two times per day.

Another preferred composition is to combine all of the above: 100mg SAMe with 60 mg Ginko Biloba Extract, and 900 mg St. John's Wort extract within a time-released coated capsule or tablet along with standard inert ingredients utilized in the art; said composition to be taken one to two times per day.

The present invention may be formulated for administration by any conventional route including but not limited to oral, rectal, nasal, topical, or pareateral. The composition may take the form of tablets, capsules, suspensions, emulsions, solutions, suppositories, sprays or injectibles, but is not limited to these forms.

Preferred Dosages

1. For the first ten days: Take 800 - 1,200 mg per day of the SAMe 400mg Active Ions tablets, caplets, or capsules - in addition to our Mind and Mood Product

2. Afterwards, continue only taking the Mind and Mood composition daily.

	WO 01/26646 PCT/US00/27559
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19	 400 mg of Active SAMe Ions 60 mg of Ginkgo Biloba Extract, 24% Flavoneglycosides - 6-7% Terpenelactones 500-900 mg of St. John's Wort Extract, .3% Hypericin & 1-2% Hyperforin. SAMe Mind and Mood Support Composition: 3. For the first ten days: Take 800 - 1,200 mg per day of our SAMe 400mg Active Ions tablets, caplets, or capsules - in addition to our Mind and Mood Product 4. Afterwards, continue only taking the Mind and Mood product daily. 400 mg of Active SAMe Ions 60 mg of Ginkgo Biloba Extract, 24% Flavoneglycosides - 6-7% Terpenelactones 500 000 mg of St. Ichn's West Extract 2% Hypericin & 1.2%
20 21 22	• 500-900 mg of St. John's Wort Extract, .3% Hypericin & 1-2% Hyperforin.
23	LIVER ENHANCEMENT AND ANTI-OXIDANT COMPOSITION
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25	The preferred liver enhancement and anti-oxidant compositions of
26	this invention preferably include a combination of SAMe and other
27	natural liver enhancement and anti-oxidant substances in effective
28	amounts for this purpose. One preferred embodiment of this invention
29	especially effective as a liver enhancement and anti-oxidant is SAMe in
30	combination with one or more of the following: Coenzyme Q-10,
31	glutathione, and frutose - 1,6 - diphosphate.
32	The following are the preferred weight percentage ranges of these
33	liver enhancement and anti-oxidant compositions:
34	
35	Component Weight Percentage
36	SAMe about 5% to about 90%

about 5% to about 90%

37

Coenzyme Q-10

	WO 01/26646				PCT/US	800/27559
1	Glutathione ab	out	5%	to	about	90%
2	Frutose - 1,6 - diphosphate ab	out	5%	to	about	90%
3						
4 5	Preferred Dosages					
6 7 8 9 10	 150 mg Co-Q-10 (Coenzyme Q-10) 400 mg SAMe 100-300 mg Glutathione 50-150 mg Frutose - 1,6 - Diphosphare 	te (F	FDP))		
11 12	SAMe Liver and Anti-Oxidant Support					
13						
14 15	150 mg Co-Q-10 (Coenzyme Q-10)400 mg SAMe					
16	• 100-300 mg Glutathione					
17	• 50-150 mg Frutose - 1,6 - Diphosphae	te (F	DP))		
18 19						
20	SLEEP ENHANCEMENT COMPO	OSIT	TOI	N		
21						
22	The preferred sleep enhancement compos					
23	preferably include a combination of SAMe					_
24	enhancement substances in effective amounts					
25	preferred embodiment of this invention especi		•			-
26	enhancement product is SAMe in combination w	vith	one	0	r more	e of the
27	following: melatonin and a Kava Kava derivative					
28	The following are the preferred weight per	cent	age	ra	inges	of these
29	sleep enhancement compositions:					
30						
31	Component	W	/eig	ht	Perce	entage
32						
33					about	
34	Melatonin ab	out :	5%	to	about	90%

about 5% to about 90%

35

Kava Kava Extract

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2	Preferred Dosages
3	
4	• 50 mg Melatonin
5	 600 mg Kava Kava Extract, 30% Kavalactones
6	• 400 mg SAMe Active Ions
7	• 100 mg Valerian Extract .8-1%
8	
9	The following are examples for the manufacture of the most preferred
10	compositions.
11	
12	Production Example I (Stress Reduction Composition)
13	1 routeron Example 1 (biress Reduction Composition)
14	Starting material:
15	
16	35.3 kilograms of Kava Kava 30% kavalactones
17	47.1 kilograms of SAMe, tosylate salt
18	5.9 kilograms of Valerian 0.8% by weight Valerenic Acid
19	200 kilograms of excepients and miscellaneous fillers
20	125,000 "00" size capsules
21	
22	• Insure that all ingredients are at approximately 2% - 3% moisture
23	content.
24	Maintain average humidity of the manufacturing environment at
25	under 25% during manufacture.
26	• Maintain average temperature of the manufacturing environment is
27	under 20 degrees Celsius.
28 29	 Make sure process time from mixture to encapsulation is under 2 hours.
30	
31	• Within 2 hours of encapsulation, enteric coat, or otherwise treat with a time release agent, the capsules for optimum stability
32	a time release agent, the capsules for optimum stability
33	
34	
35	Production Example II (Mind and Mood Enhancement
36	Reduction Composition)
37	
38	Starting material:
39	
40	50 kilograms of SAMe, tosylate salt
41	4 kilograms of Ginkgo Biloba Extract 24/6% by weight Extract
42	31 kilograms of St. John's Wort, 3% by weight Extract
43	240 kilograms of excepients and miscellaneous fillers
44	125,000 "00" size capsules

WO 01/26646 PCT/US00/27559 1 2 • Insure that all ingredients are at approximately 2% - 3% moisture 3 content. 4 • Maintain average humidity of the manufacturing environment is at 5 • Maintain average temperature of the manufacturing environment is 6 7 under 20 degrees Celsius 8 • Make sure process time from mixture to encapsulation is under 2 9 hours. • Within 2 hours of encapsulation, enteric coat, or otherwise treat with 10 a time release agent, the capsules for optimum stability 11 12 13 Production Example III (Joint Anti-Inflammation Composition) 14 15 16 Starting material: 17 18 33.4 kilograms of SAMe, tosylate salt 19 41.7 kilograms glucosamine sulfate 20 12.5 kilograms of methyl sulfanyl methane (MSN) 21 kilograms of a 20% by weight cetyl myristoleate blend. 22 10.4 kilograms of excepients and miscellaneous fillers 125,000 "00" size capsules 23 24 25 • Insure that all ingredients are at approximately 2% - 3% moisture 26 content. • Maintain average humidity of the manufacturing environment is 27 28 under 25% • Maintain average temperature of the manufacturing environment is 29 at under 20 degrees Celsius 30 • Make sure process time from mixture to encapsulation is under 2 31 32 hours • Within 2 hours of encapsulation, enteric coat, or otherwise treat with 33 a time release agent, the capsules for optimum stability 34 35 36 Production Example IV (Weight Reduction and Energy 37 Enhancement Composition) 38 39 Starting material: 40 41

50 kilograms of SAMe, tosylate salt

19 kilograms Ma Huang Extract 6% by weight

6.3 kilograms of Epimedium Pinnatum Extract 10% by weight

6.3 kilograms of Citrus Aurantium Extract 6% by weight

5 kilograms of Guarana Extract 10% by weight

42 43

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	WO 01/26646 PCT/US00/27559
1	13.7 kilograms of excepients and miscellaneous fillers
2	125,000 "00" size capsules
3	•
4	• Insure that all ingredients are at approximately 2% - 3% moisture
5	content.
6	• Maintain average humidity of the manufacturing environment is at
7	under 25%.
8 9	 Maintain average temperature of the manufacturing environment is under 20 degrees Celsius.
10	• Make sure process time from mixture to encapsulation is under 2
11	hours.
12	• Within 2 hours of encapsulation, enteric coat, or otherwise treat with
13	a time release agent, the capsules for optimum stability
14	a time release agent, the capsules for optimum stability
15	
16	
17	Production Example V (Liver Enhancement and Anti-Oxident
	Production Example V (Liver Enhancement and Anti-Oxidant
18	Composition)
19	Canting materials
20	Starting material:
21	05.0.111
22	25.8 kilograms of SAMe, tosylate salt
23	9.7 kilograms Co-Q-10
24	6.5 kilograms glutahion
25	6.4 kilograms fructose – 1,6 – diphosphate
26	25.8 kilograms of excepients and miscellaneous fillers
27	125,000 "00" size capsules
28	
29	• Insure that all ingredients are at approximately 2% - 3% moisture
30	content.
31	• Maintain average humidity of the manufacturing environment is at
32	under 25%.
33	• Maintain average temperature of the manufacturing environment is
34	under 20 degrees Celsius.
35	• Make sure process time from mixture to encapsulation is under 2
36	hours.
37	• Within 2 hours of encapsulation, enteric coat, or otherwise treat with
38	a time release agent, the capsules for optimum stability
39	estimate and the second
40	
41	Production Example VI (Sleep Enhancement Composition)
42	, and the same of
43	Starting material:
44	
45	47.1 kilograms of SAMe, tosylate salt
_	

1	3.0 kilograms of melatonin
2	35.5 kilograms of Kava Kava Extract
3	5.9 kilograms of Valerian Root Extract
4	8.9 kilograms of excepients and misc. fillers
5	125,000 "00" size capsules

6

- Insure that all ingredients are at approximately 2% 3% moisture content.
- 9 Maintain average humidity of the manufacturing environment is at under 25%.
- Maintain average temperature of the manufacturing environment is under 20 degrees Celsius.
- Make sure process time from mixture to encapsulation is under 2 hours.
- Within 2 hours of encapsulation, enteric coat, or otherwise treat with a time release agent, the capsules for optimum stability

17 18

SCOPE OF THE INVENTION

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The above presents a description of the best mode contemplated of carrying out the present invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains to make and this invention. This invention is, however, susceptible modifications and alternate constructions from that discussed above which are fully equivalent. Consequently, it is not the intention to limit this invention to the particular embodiments disclosed. On the contrary, the intention is to cover all modifications and alternate constructions coming within the spirit and scope of the invention as generally expressed by the following claims, which particularly point out and distinctly claim the subject matter of the invention:

1 THE CLAIMS

2

- 3 1. A nutraceutical product comprising a mixture of SAMe and Co-Q-
- 4 10.

5

- 6 2. The nutraceutical product of Claim 1 comprising from 0.1 to 99
- 7 weight percent of the SAMe and from 1 to 99 weight percent of the Co-
- 8 Q-10.

9

- 10 3. The nutraceutical product of Claim 2 where the moisture content
- 11 of the product is less than 5 weight percent.

12

- 13 4. A nutraceutical product comprising a mixture SAMe and St. John's
- 14 Wort derivative.

15

- 16 5. The nutraceutical product of Claim 4 comprising from 0.1 to 99
- 17 weight percent of the SAMe and from 1 to 99 weight percent of the St.
- 18 John's Wort derivative.

19

- 20 6. The nutraceutical product of Claim 5 where the moisture content
- 21 of the product is less than 5 weight percent.

22

- 23 7. A nutraceutical product comprising a mixture SAMe and a Ginkgo
- 24 Biloba derivative.

25

- 26 8. The nutraceutical product of Claim 7 comprising from 0.1 to 99
- 27 weight percent of the SAMe and from 1 to 99 weight percent of the
- 28 Ginkgo Biloba derivative.

1 9. The nutraceutical product of Claim 8 where the moisture content

2 of the product is less than 5 weight percent.

3

- 4 10. A nutraceutical product comprising a mixture SAMe and a Ma
- 5 Huang derivative.

6

- 7 11. The nutraceutical product of Claim 10 comprising from 0.1 to 99
- 8 weight percent of the SAMe and from 1 to 99 weight percent of the Ma
- 9 Huang derivative.

10

- 11 12. The nutraceutical product of Claim 11 where the moisture content
- of the product is less than 5 weight percent.

13

- 14 13. A nutraceutical product comprising a mixture SAMe and a
- 15 Epimeduim Pinnatum derivative.

16

- 17 14. The nutraceutical product of Claim 13 comprising from 0.1 to 99
- 18 weight percent of the SAMe and from 1 to 99 weight percent of the
- 19 Epimeduim Pinnatum derivative.

20

- 21 15. The nutraceutical product of Claim 14 where the moisture content
- of the product is less than 5 weight percent.

23

- 24 16. A nutraceutical product comprising a mixture SAMe and a
- 25 Guarana derivative.

26

- 27 17. The nutraceutical product of Claim 16 comprising from 0.1 to 99
- 28 weight percent of the SAMe and from 1 to 99 weight percent of the
- 29 Guarana derivative.

1 18. The nutraceutical product of Claim 17 where the moisture content

2 of the product is less than 5 weight percent.

3

- 4 19. A nutraceutical product comprising a mixture SAMe and a Citrus
- 5 Aurantium derivative.

6

- 7 20. The nutraceutical product of Claim 19 comprising from 0.1 to 99
- 8 weight percent of the SAMe and from 1 to 99 weight percent of the
- 9 Citrus Aurantium derivative.

10

- 11 21. The nutraceutical product of Claim 20 where the moisture content
- 12 of the product is less than 5 weight percent.

13

- 14 22. A nutraceutical product comprising a mixture SAMe and a Kava
- 15 Kava derivative.

16

- 17 23. The nutraceutical product of Claim 22 comprising from 0.1 to 99
- 18 weight percent of the SAMe and from 1 to 99 weight percent of the
- 19 Kava Kava derivative.

20

- 21 24. The nutraceutical product of Claim 23 where the moisture content
- 22 of the product is less than 5 weight percent.

23

- 24 25. A nutraceutical product comprising a mixture SAMe and cetyl
- 25 myristoleate.

26

- 27 26. The nutraceutical product of Claim 25 comprising from 0.1 to 99
- 28 weight percent of the SAMe and from 1 to 99 weight percent of the
- 29 cetyl myristoleate.

1 27. The nutraceutical product of Claim 29 where the moisture content

2 of the product is less than 5 weight percent.

3

- 4 28. The nutraceutical product of Claim 27 including cetyl oleate, cetyl
- 5 esters, or cetyl myristate.

6

- 7 29. The nutraceutical product of Claim 28 comprising from 20 to 80
- 8 weight percent of the cetyl myristoleate and comprising from 20 to 80
- 9 weight percent of the cetyl oleate, cetyl esters, or cetyl myristate.

10

- 11 30. A nutraceutical product comprising a mixture SAMe and a
- 12 Valerian derivative.

13

- 14 31. The nutraceutical product of Claim 33 comprising from 0.1 to 99
- 15 weight percent of the SAMe and from 1 to 99 weight percent of the
- 16 Valerian derivative.

17

- 18 32. The nutraceutical product of Claim 31 where the moisture content
- 19 of the product is less than 5 weight percent.

20

- 21 33. A nutraceutical product comprising a mixture SAMe and
- 22 glutathione.

23

- 24 34. The nutraceutical product of Claim 33 comprising from 0.1 to 99
- 25 weight percent of the SAMe and from 1 to 99 weight percent of the
- 26 glutathione.

27

- 28 35. The nutraceutical product of Claim 34 where the moisture content
- 29 of the product is less than 5 weight percent.

1 36. A nutraceutical product comprising a mixture SAMe and fructose -

2 1,6-diphosphate.

3

- 4 37. The nutraceutical product of Claim 36 comprising from 0.1 to 99
- 5 weight percent of the SAMe and from 1 to 99 weight percent of the
- 6 fructose 1,6 diphosphate.

7

- 8 38. The nutraceutical product of Claim 37 where the moisture content
- 9 of the product is less than 5 weight percent.

10

- 11 39. A nutraceutical product comprising a mixture SAMe and
- 12 glucosamine sulfate, glucosamine HCL or glucosamine potassium.

13

- 14 40. The nutraceutical product of Claim 39 comprising from 0.1 to 99
- 15 weight percent of the SAMe and from 1 to 99 weight percent of the
- 16 glucosamine sulfate, HCL or potassium.

17

- 18 41. The nutraceutical product of Claim 40 where the moisture content
- 19 of the product is less than 5 weight percent.

20

- 21 42. A nutraceutical product comprising a mixture SAMe and methyl
- 22 sulfanyl methane.

23

- 24 43. The nutraceutical product of Claim 42 comprising from 0.1 to 99
- 25 weight percent of the SAMe and from 1 to 99 weight percent of the
- 26 methyl sulfanyl methane.

27

- 28 44. The nutraceutical product of Claim 43 where the moisture content
- of the product is less than 5 weight percent.

A nutraceutical product comprising a mixture SAMe and a dietary 1

- supplement, with the moisture content of the product being less than 5 2
- 3 weight percent.

4

- 5 The nutraceutical product of Claim 45 where the SAMe is a
- mixture of (RS)-(+)-SAMe and (SS)-(+)-SAMe diastereoisomers, with the 6
- 7 (SS)-(+)-SAMe diastereoisomer being at a concentration of at least 95
- 8 weight percent of the mixture.

9

- 10 A nutraceutical product comprising a softgel pill, caplet, capsule
- 11 or tablet treated with a time release agent, said softgel pill, caplet,
- 12 capsule or tablet containing a mixture SAMe and a dietary supplement,
- with the moisture content of the mixture being less than 5 weight 13
- 14 percent.

15

- The nutraceutical product of Claim 47 where the softgel pill, 16 48.
- 17 caplet, capsule or table sealed within an aluminum foil package that acts
- 18 as a moisture barrier, enabling said softgel pill, caplet, capsule or tablet
- to have substantially prolonged life when subjected to accelerated 19
- 20 testing than a softgel pill, caplet, capsule or table contained within a
- 21 sealed bottle.

22

- 23 The nutraceutical product of Claim 47 where the time release
- 24 agent is an aqueous ethyl cellulose dispersion.

25

- 26 The nutraceutical product of Claim 47 where the SAMe is a 50.
- mixture of (RS)-(+)-SAMe and (SS)-(+)-SAMe diastereoisomers, with the 27
- 28 (SS)-(+)-SAMe diastereoisomer being at a concentration of at least 95
- 29 weight percent of the mixture.

1 51. A nutraceutical product comprising a mixture of SAMe, a Kava

2 Kava derivative, and a Valerian derivative.

3

- 4 52. The nutraceutical product of Claim 51 where the Kava Kava
- 5 derivative includes from 30 to 50 weight percent Kavalactones.

6

- 7 53. The nutraceutical product of Claim 51 where the Valerian
- 8 derivative includes from 0.8 to 1 weight percent Valerenic acid.

9

- 10 54. A nutraceutical product comprising a mixture of SAMe, a Ginkgo
- 11 Biloba derivative, and a St. John's Wort derivative.

12

- 13 54. A nutraceutical product comprising a mixture of SAMe,
- 14 glucosamine suffate, methyl sulfanyl methane, and cetyl myristoleate.

15

- 16 55. A nutraceutical product comprising a mixture of SAMe, Ma Huang
- 17 derivative, Epimedium Pinnatum, derivative, Citrus Aurantium
- 18 derivative, Guarana derivative.

19

- 20 56. A nutraceutical product comprising a mixture of SAMe, Co-Q-10,
- 21 glutahion, and fructose 1,6 diphosphate.

22

- 23 57. A nutraceutical product comprising a mixture of SAMe, a Kava
- 24 Kava derivative, a Valerian derivative, and melatonin.

- 26 58. A nutraceutical product comprising a mixture of SAMe and at least
- 27 two of the following ingredients:
- 28 Co-Q-10 (Coenzyme Q-10),
- 29 St. John's Wort derivative, either powder or any form of extract,
- Ginkgo Biloba derivative, either powder or any form of extract,

- 1 Ma Huang derivative, either powder or any form of extract,
- 2 Epimeduim pinnatum derivative, either powder or extract,
- 3 Guarana derivative, either leaf or root powder or any form of
- 4 extract,
- 5 Citrus Aurantium derivative, either powder or any from of extract,
- 6 Kava Kava derivative, either powder or any form of extract,
- 7 cetyl myristoleate (CMO),
- 8 cetyl myristoleate blended with cetyl oleate, cetyl esters and cetyl
- 9 myristate,
- Valerian derivative, either powder or any form of extract,
- 11 glutathione,
- fructose 1,6 diphosphate (FDP),
- glucosamine sulfate, glucosamine HCL, or glucosamine potassium,
- methyl sulfanyl methane (MSM),
- 15 melatonin,
- 16 hydrolyzed collagen,
- 17 chondroitin sulfate,
- 18 citicoline,
- 19 alpha GPC.

20

- 21 57. A nutraceutical product comprising a mixture of SAMe and
- 22 melatonin.

23

- 24 58. A nutraceutical product comprising a mixture of SAMe and
- 25 hydrolyzed collagen.

- 27 59. A nutraceutical product comprising a mixture of SAMe and
- 28 chondroitin sulfate,
- 29 60. A nutraceutical product comprising a mixture of SAMe and
- 30 citicoline,

1 61. A nutraceutical product comprising a mixture of SAMe and alpha

2 GPC.

3

- 4 62. A method of manufacturing a nutraceutical product comprising
- 5 mixing SAMe and a dietary supplement under conditions where the
- 6 moisture content of the product is less than 5 weight percent.

7

- 8 63. The method of Claim 62 where the humidity of the manufacturing
- 9 environment of nutraceutical product is less than 25%.

10

- 11 64. The method of Claim 62 where the temperature of the
- 12 manufacturing environment of nutraceutical product is from 20 to 15
- 13 degrees Celsius.

14

- 15 65. The method of Claim 64 where the nutraceutical product is
- 16 encapsulated with capsules within 2 hours after manufacture.

17

- 18 66. The method of Claim 65 where the capsules are enteric coat, or
- 19 otherwise treat with a time release agent, within 2 hours after
- 20 encapsulation.

21

- 22 67. A method of supplementing a diet by administering for at least 10
- 23 to 14 days SAMe at a dosage of from 800 to 1200 milligrams per day
- 24 and at the same time administering a nutraceutical product comprising
- 25 SAMe and a dietary supplement, said nutraceutical product including a
- 26 a dosage of at least 400 milligrams per day of SAMe, and after 14 days
- 27 administering only said nutraceutical product.

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INTERNATIONAL SEARCH REPORT

International application No. PCT/US00/27559

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	to International Patent Classification (IPC) or to both	national classification and IPC			
B. FIEL	DS SEARCHED				
Minimum d	ocumentation searched (classification system followers	ed by classification symbols)			
	424/195.1, 514/ 46, 125, 415, 560, 689	,,			
0.5	727/1/3:1, 314/ 10, 123, 113, 300, 003				
Documenta	tion searched other than minimum documentation to th	e extent that such documents are included	in the fields searched		
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Electronic d	lata base consulted during the international search (n	ame of data base and, where practicable	, search terms used)		
REGISTR	Y, CAPLUS, NAPRALERT, BIOSIS, MEDLINE,	EMBASE, WPIDS			
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C. DOC	UMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where a	ppropriate, of the relevant passages	Relevant to claim No.		
Y	Database CAPLUS on STN (Col	lumbus, OH, USA), No.	1-67		
	129:347310, HENDERSON ET				
	glycosaminoglycan or glycosaminogly				
	adenosylmethionine,' abstract, WO 98	48816 A1, 19981105.			
Y	Database CAPLUS on STN, (col		1-67		
	130:329206, HENRIKSEN, B 'Pha				
	treating liver disorders containin inorg	ganic selenium and vitamins,			
	abstract, EP 913155 A2, 19990506.				
		i	;		
Furth	er documents are listed in the continuation of Box C	See patent family annex.			
• Spi	ecial categories of cited documents:	"T" later document published after the inte date and not in conflict with the appli	rnational filing date or priority		
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- (71) Applicant (for all designated States except US): PHARMNSEAS, INC. [US/US]; 22512 Avenida Empresa, Rancho Santa Margarita, CA 92688 (US).
- (72) Inventor; and
- (75) Inventor/Applicant (for US only): HOWARD, Larry [US/US]; 25262 Arcadian Avenue, Mission Viejo, CA 92691 (US).
- (74) Agent: CONNORS, John, J.; Connors & Associates, Suite 220, 1600 Dove Street, Newport Beach, CA 92660-2427 (US).

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(54) Title: NUTRACEUTICAL PRODUCTS CONTAINING SAME AND DIETARY SUPPLEMENTS AND METHOD OF MAN-UFACTURING AND USE THEREOF

(57) Abstract: A nutraceutical product comprises a mixture SAMe and a dietary supplement, where the moisture content of the product is less than 5 weight percent. A nutraceutical preferred product includes a mixture of (RS)-(+)-SAMe and (SS)-(+)-SAMe diastereoisomers, with the (SS)-(+)-SAMe diastereoisomer being at a concentration of at least 95 weight percent of the mixture.